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<p>(21) International Application Number: PCT/DK89/00096 (22) International Filing Date: 25 April 1989 (25.04.89) (30) Priority data: 2254/88 26 April 1988 (26.04.88) DK (71) Applicant (for all designated States except US): NOVO-NORDISK A/S [DK/DK]; Novo Alle, DK-2880 Bagsværd (DK). (72) Inventors; and: (75) Inventors/Applicants (for US only): NORRIS, Kjeld [DK/DK]; Ahlmanns Alle 34, DK-2900 Hellerup (DK). PETERSEN, Lars, Christian [DK/DK]; Havrevej 4, DK-2960 Hørsholm (DK). (74) Common Representative: NOVO-NORDISK A/S; Patent Department, Novo Alle, DK-2880 Bagsværd (DK).</p>		<p>(81) Designated States: AU, DK, FI, JP, NO, US.  <b>Published</b> <i>With international search report.</i></p>
<p>(54) Title: APROTININ ANALOGUES AND A PROCESS FOR THE PRODUCTION THEREOF</p> <pre> 3       5       10      15 AspPheCysLeuGluProProTyrThrGlyProCysLysAlaArgIle AAAGAGATTTCTGTTTGAACCTCCATACACTGGTCCATGTAAAGCTAGAATC CTAAAGACAAACCTTGGAGGTATGTGACCAGGTACATTTCGATCTTAG PflMI  20      25      30      35 IleArgTyrPheTyrAsnAlaLysAlaGlyLeuCysGlnThrPheValTyrGly ATCAGATACTTCTACAACGCAAGGCTGGTTGTGTCAAACCTTTCGTTTACGGT TAGTCTATGAAGATGTTGCGGTTCCGACCAAACACAGTTTGAAAGCAAATGCCA StyI  40      45      50 GlyCysArgAlaLysSerAsnAsnPheLysSerAlaGluAspCysMetArgThr GGCTGCAGAGCTAAGTCCAACAACCTCAAGTCTGCTGAAGACTGCATGAGAACT CCGACGTCTCGATTACAGTTGTTGAAGTTCAGACGACTTCTGACGTACTCTTGA PstI  55      58 CysGlyGlyAlaStop TGTGGTGGTGCCTAAT ACACCACCACGGATTAGATC XbaI </pre> <p>(57) Abstract</p> <p>Novel aprotinin analogues having a selected inhibition profile against serine proteases.</p>		

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Aprotinin Analogues and a Process for the Production Thereof

## FIELD OF THE INVENTION

The present invention relates to novel aprotinin analogues and to a process for their production.

5

## BACKGROUND OF THE INVENTION

Throughout the present specification the term  
10 naturally occurring amino acid (or amino acid residue)  
refers to one of the  $\alpha$ -amino acids which are listed below  
together with the symbols used to designate the individual  
amino acid residues:

15

	Asp	Aspartic acid	Ile	Isoleucine
	Thr	Threonine	Leu	Leucine
	Ser	Serine	Tyr	Tyrosine
	Glu	Glutamine acid	Phe	Phenylalanine
20	Pro	Proline	His	Histidine
	Gly	Glycine	Lys	Lysine
	Ala	Alanine	Arg	Arginine
	Cys	Cysteine	Trp	Tryptophane
	Val	Valine	Gln	Glutamine
25	Met	Methionine	Asn	Asparagine

All amino acids (or amino acid residues)  
mentioned in the present specification have the L-  
30 configuration, except glycine which has no chiral center.

Aprotinin (bovine pancreatic trypsin inhibitor, BPTI) is a polypeptide present in several bovine organs and tissues, such as lymph nodes, pancreas, lung, parotid gland, spleen and liver. It is a single chain polypeptide consisting of 58 amino acid residues cross-linked by three disulphide bridges in the following sequence:

```

Arg-Pro-Asp-Phe-Cys-Leu-Glu-Pro-Pro-Tyr-Thr-Gly-Pro-Cys-
1  2  3  4  5  6  7  8  9  10 11 12 13 14
10: Lys-Ala-Arg-Ile-Ile-Arg-Tyr-Phe-Tyr-Asn-Ala-Lys-Ala-Gly-
    15 16 17 18 19 20 21 22 23 24 25 26 27 28
    Leu-Cys-Gln-Thr-Phe-Val-Tyr-Gly-Gly-Cys-Arg-Ala-Lys-Arg-
    29 30 31 32 33 34 35 36 37 38 39 40 41 42
    Asn-Asn-Phe-Lys-Ser-Ala-Glu-Asp-Cys-Met-Arg-Thr-Cys-Gly-
15  43 44 45 46 47 48 49 50 51 52 53 54 55 56
    Gly-Ala
    57 58

```

The three disulphide bridges are situated between Cys(5)-Cys(55), Cys(14)-Cys(38) and Cys(30)-Cys(51), respectively.

Aprotinin inhibits various serine proteases, such as trypsin, chymotrypsin, plasmin and kallikrein and is used for the treatment of acute pancreatitis, various states of shock syndroms, hyperfibrinolytic haemorrhage, and myocardial infarction. Administration of aprotinin in high doses significantly reduces blood loss in connection with cardiac surgery.

Aprotinin is extracted from various bovine organs or tissues, such as lung, pancreas and parotid glands. Extraction from animal tissues is a cumbersome process and requires large amounts of the bovine organ or tissue. Aprotinin may also be produced by recombinant DNA-technology by insertion of a gene coding for aprotinin in a suitable microorganism which when cultured in a suitable nutrient medium produces the desired product.

Production of aprotinin analogues in E. coli is described in EP published patent application No. 238,993 and production of aprotinin in yeast is described in Danish patent application No. 4501/87.

5 Certain aprotinin analogues and derivatives have been described, see for instance Jering H. and Tschesche H., Eur.J.Biochem. 61 (1976), 453-463 describing replacement of Lys(15) with Arg, Phe or Trp or US patent specification No. 4,595,674 describing aprotinin analogues  
10 in which the lysine residue in position 15 in the active center of the aprotinin has been replaced by Gly, Ala, Val, Leu, Ile, Met, Arg, L- $\alpha$ -amino butyric acid, L-norvaline, L-norleucine, dehydroalanine or L-homoserine. Also, the above mentioned EP No. 238,993 describes aprotinin  
15 analogues having Lys(15) substituted by Arg, Val, Ile, Leu, Phe, Gly, Ser, Trp, Tyr or Ala and/or Met(52) substituted by Glu, Leu, Val, Thr or Ser.

The known aprotinin analogues are claimed to have modified effects and efficacies towards different  
20 proteinases. For instance aprotinin(15Val) has a relatively high selectivity for granulocyte elastase and an inhibition effect on collagenase, aprotinin(15Ala) has only a weak inhibitory effect on elastase and aprotinin(15Gly) has an outstanding antitrypsin activity and surprisingly inhibits  
25 kallikrein.

## SUMMARY OF THE INVENTION

Is is the purpose of the present invention to provide novel aprotinin analogues having a more specific  
5 inhibitory effect towards certain serine proteases, such as elastase, kallikrein, t-PA, urokinase and coagulation factors, such as thrombin.

The present invention is based on the surprising fact that replacement of Arg in position 17 of  
100 aprotinin(3-58;42Ser) with Ala or of Arg in position 17 with Ala and of Ile in position 19 of aprotinin (3-58;42Ser) with Glu gives rise to a substantial increase in the inhibition of human plasma kallikrein. This is even more pronounced on replacement of Lys in position 15 with  
15 Arg and of Arg in position 17 with Ala in aprotinin (3-58;42Ser).

Accordingly, the present invention is related to aprotinin analogues in which at least one of the amino acid residues 16 to 19 have been replaced by another naturally  
20 occurring amino acid residue.

In the above mentioned Danish patent application No. 4501/87 aprotinin analogues with certain amino acid residue substitutions and/or deletions are described. The aim of these amino acid residue substitutions and/or  
25 deletions is to avoid proteolytical cleavage at certain basic amino acid residues during the production of the aprotinin analogue in yeast. In particular amino acid residues 1 and 2 may be deleted and one of the basic amino acid residues at the dibasic sequence  
30 41 - 42 may be replaced by another amino acid residue. It has been shown that such aprotinin analogues are produced in high yields in yeast and exhibit the same characteristics as native aprotinin.

To ensure high production in yeast the present aprotinin analogues may further be modified in a similar way. Accordingly the present aprotinin analogues besides being modified in the sequence from amino acid 16 to 19 may  
5 also be modified at sequence 1 - 2 and 41 - 42, respectively provided that such further modification does not have an adverse effect on the goal of the present invention.

The present aprotinin analogues may furthermore  
10 be modified in position 15 by insertion at this position of another naturally occurring amino acid residue including the known substitutions with naturally occurring amino acid residues.

Native aprotinin contains a disulphide bridge  
15 between Cys(14) and Cys(38) which is strongly involved in the tertiary structuring around the active site at Lys(15). This disulphide bond may be split by reducing agents. A more convenient way to avoid having a disulphide bridge at this position in the molecule would be to substitute the  
20 cysteine residues with other residues or simply to delete these residues. Accordingly, the present aprotinin analogues may furthermore be modified so that they do not contain a disulphide bridge between Cys(14) and Cys(38). This might impart further interesting characteristics to  
25 the molecule.

Finally modifications at positions 12 and 13 might be considered.

Accordingly in its broadest aspect the present invention relates to aprotinin analogues being modified in  
30 the sequence from amino acid residue 12 to 19 provided that if Lys(15) has been replaced by another amino acid then at least one further amino acid residue in said sequence has

been substituted or deleted. The present aprotinin analogues may furthermore as described above be modified in the known way at the sequences 1 - 2 and 41 - 42. . .

According to a further aspect of the present  
5 invention there is provided a method for producing the  
above aprotinin analogues by cultivation of a yeast strain  
containing a replicable expression vector containing a gene  
encoding the aprotinin analogues in a suitable nutrient  
medium followed by recovery of the desired product from the  
10 culture medium.

#### BREIF DESCRIPTION OF THE DRAWINGS

15 The present invention will be further  
illustrated by reference to the accompanying drawings in  
which:

Fig. 1 shows a synthetic gene encoding aprotinin  
(3-58; 42Ser);

20

Fig. 2 illustrates the construction of the plasmid  
pKFN 306;

25

Fig. 3 illustrates the construction of the plasmid  
pKFN 504;

Fig. 4 illustrates the construction of the plasmid  
pMT 636;

30

Fig. 5A illustrates the inhibition of plasma  
kallikrein by native aprotinin and by the  
aprotinin analogues KFN 396 and KFN 399; and



Fig. 5B illustrates the inhibition of plasma kallikrein by native aprotinin and by the aprotinin analogues KFN 396, KFN 772 and KFN 773.

5

#### DETAILED DESCRIPTION OF THE PRESENTLY PREFERRED EMBODIMENTS

The present aprotinin analogues may be represented by the following formula (I):

10.

$X_1$ -Asp-Phe-Cys-Leu-Glu-Pro-Pro-Tyr-Thr- $X_2$ - $X_3$ - $X_4$ - $X_5$ - $X_6$ - $X_7$ -  
 $X_8$ - $X_9$ -Arg-Tyr-Phe-Tyr-Asn-Ala-Lys-Ala-Gly-Leu-Cys-Gln-Thr-  
 Phe-Val-Tyr-Gly-Gly- $X_{10}$ -Arg-Ala- $X_{11}$ - $X_{12}$ -Asn-Asn-Phe-Lys-  
 Ser-Ala-Glu-Asp-Cys-Met-Arg-Thr-Cys-Gly-Gly-Ala

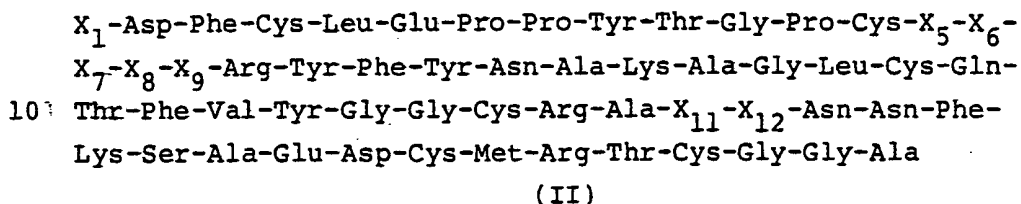
15

(I)

in which  $X_1$  is Arg-Pro, Pro or hydrogen, preferably Arg-Pro or hydrogen, most preferred hydrogen;  $X_2$  is any naturally occurring amino acid residue, preferably Gly;  $X_3$  is any  
 20 naturally occurring amino acid residue, preferably Pro;  $X_4$  and  $X_{10}$  are each any naturally occurring amino acid residue, preferably they are both Cys or both Ala, most preferred they are both Cys;  $X_5$  is Lys, Arg, Val, Thr, Ile, Leu, Phe, Gly, Ser, Met, Trp, Tyr or Ala, preferably Lys or Arg, with  
 25 the proviso that when  $X_5$  is different from Lys, then at least one of  $X_2$  to  $X_4$  and  $X_6$  to  $X_9$  are different from the amino acid residue at the corresponding position in native aprotinin;  $X_6$  is Ala or Gly, preferably Ala;  $X_7$  is any naturally occurring amino acid residue, preferably Ala or  
 30 Gly;  $X_8$  is Ile, Leu, Met, Val or Phe, preferably Ile;  $X_9$  is any naturally occurring amino acid residue, preferably Ile or Glu;  $X_{11}$  is any naturally occurring amino acid residue, preferably Lys or Arg;  $X_{12}$  is Lys, Arg or Ser at least one

of the amino acid residues  $X_2$  to  $X_9$ , preferably  $X_5$  to  $X_9$ , more preferred  $X_6$  to  $X_9$  being different from the corresponding amino acid residue in native aprotinin.

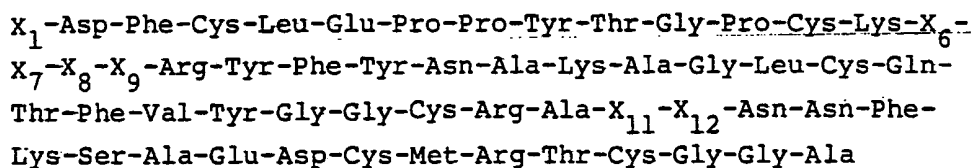
According to a more narrow aspect the present  
5 aprotinin analogues may be represented by the following formula (II):



in which  $X_1$ ,  $X_5$ ,  $X_6$ ,  $X_7$ ,  $X_8$ ,  $X_9$ ,  $X_{11}$  and  $X_{12}$  are as defined  
above for formula (I), at least one of the amino acid  
15 residues  $X_5$  to  $X_9$ , preferably  $X_6$  to  $X_9$  being different from  
the corresponding amino acid residue in native aprotinin.

According to an even narrower aspect the  
aprotinin analogues may be represented by the following  
formula (III):

20



25

(III)

in which  $X_1$ ,  $X_6$ ,  $X_7$ ,  $X_8$ ,  $X_9$ ,  $X_{11}$  and  $X_{12}$  are as defined  
above for formula (I), at least one of the amino acid  
residues  $X_6$  to  $X_9$  being different from the corresponding  
amino acid residue in native aprotinin.

30

Examples of preferred aprotinin analogues  
according to the present invention are aprotinin(3-58;  
17Ala + 42Ser), which lacks the first two amino acid  
residues of native aprotinin and has Ala substituted for

Arg in position 17 and Ser substituted for Arg in position 42; aprotinin (3-58; 17Ala + 19Glu + 42Ser), which lacks the first two amino acid residues of native aprotinin and has Ala substituted for Arg in position 17, Glu substituted  
5 for Ile in position 19 and Ser substituted for Arg in position 42; and aprotinin (3-58; 15Arg + 17Ala + 42Ser) which lacks the first two amino acid residues of native aprotinin and has Arg substituted for Lys in position 15, Ala substituted for Arg in position 17 and Ser substituted  
10 for Arg in position 42, respectively.

Further examples of aprotinin analogues according to the present invention are:

15 Aprotinin(3-58; 17Ala)  
Aprotinin(3-58; 17Ala + 19Glu)  
Aprotinin(3-58; 15Arg + 17Ala)  
Aprotinin(17Ala + 42Ser)  
Aprotinin(15Arg + 17Ala + 42Ser)  
Aprotinin(17Ala)  
20 Aprotinin(17Ala + 19Glu)  
Aprotinin(15Arg + 17Ala)

For secretion purposes the DNA-sequence encoding the desired aprotinin analogue is fused to a DNA-sequence  
25 encoding a signal and leader peptide sequence. The signal and leader peptides are cleaved off by the transformed microorganism during the secretion of the expressed protein product from the cells ensuring a more simple isolation procedure of the desired product. A well suited leader  
30 peptide system for yeast is the yeast MF $\alpha$ 1 leader sequence or a part thereof (Kurjan, J. and Herskowitz, I., Cell 30 (1982) 933-943). However, any signal- or leader-sequence

which provides for secretion in yeast may be employed and the present invention is not contemplated to be restricted to a specific secretion system.

For expression purposes a promoter sequence is positioned upstream to the DNA-sequence for the desired protein product. Preferably a promoter from a gene indigenous to the yeast host organism is used, e.g. the promoter of the TPI-(triose phosphate isomerase) gene. The DNA-sequence for the desired product will be followed by a transcription terminator sequence, preferably a terminator sequence from a gene indigenous to the host yeast organism, e.g. the terminator of the TPI-gene or the MF $\alpha$ 1 gene.

The DNA-sequence encoding the desired aprotinin analogue fused to appropriate promoter, signal, leader and terminator sequences is inserted into an expression vector for expression of the aprotinin analogue in yeast.

The expression vector may be a plasmid capable of independent replication in yeast or capable of integration into the yeast chromosome. The plasmid may preferably be stabilized against plasmid loss by the host microorganism by incorporation of a gene essential for the viability or normal growth of the host cells, e.g. a gene coding for cell division, cell wall biosynthesis, protein synthesis, etc.

25

#### Example 1

##### Aprotinin(3-58; 17Ala + 42Ser)(KFN396)

30

A sequence encoding aprotinin(3-58; 42Ser) was constructed from a number of oligonucleotides by ligation.

The oligonucleotides were synthesized on an automatic DNA synthesizer using phosphoramidite chemistry on a controlled pore glass support (S.L. Beaucage and M.H. Caruthers (1981) Tetrahedron Letters 22, 1859-1869).

5

The following 10 oligonucleotides were synthesized:

- |    |  |            |
|----|--|------------|
| 10 | I: AAAGAGATTCTGTTTGGAACTCCATACACTGGTCC       | } Duplex A |
|    | 37-mer                                       |            |
|    | II: TTACATGGACCAGTGTATGGAGGTTCCAAACAGAACT    | }          |
|    | 38-mer                                       |            |
| 15 | III: ATGTAAAGCTAGAATCATCAGATACTTCTACAACG     | } Duplex B |
|    | 35-mer                                       |            |
|    | IV: CTTGGCGTTGTAGAAGTATCTGATGATTCTAGCT       | }          |
|    | 34-mer                                       |            |
| 20 | V: CCAAGGCTGGTTTGTGTCAAAC TTTCGTTTACGGTGGCT  | } Duplex C |
|    | 39-mer                                       |            |
|    | VI: CTCTGCAGCCACCGTAAACGAAAGTTTGACACAAACCAGC | }          |
| 25 | 40-mer                                       |            |
|    | VII: GCAGAGCTAAGTCCAACAACTTCAAGT             | } Duplex D |
|    | 27-mer                                       |            |
| 30 | VIII: AGCAGACTTGAAGTTGTTGGACTTAG             | }          |
|    | 26-mer                                       |            |

IX:	CTGCTGAAGACTGCATGAGAACTTGTGGTGGTGCCTAAT	}	Duplex E
	39-mer		
X:	CTAGATTAGGCACCACCACAAGTTCTCATGCAGTCTTC		
5	38-mer		

5 duplexes A - E were formed from the above 10 oligonucleotides as shown in fig. 1 and 2.

10 20 pmole of each of the duplexes A - E was formed from the corresponding pairs of 5'-phosphorylated oligonucleotides I - X by heating for 5 min. at 90°C followed by cooling to room temperature over a period of 75 minutes. The five duplexes were mixed and treated with T4  
15 ligase. The synthetic sequence was isolated as a 176 bp band after electrophoresis of the ligation mixture on a 2% agarose gel.

The synthetic sequence was ligated to a 330 bp EcoRI-HgaI fragment from plasmid pKFN9 coding for MFl signal and leader sequence(1-85) and to the large EcoRI-XbaI fragment from pUC19. The construction of pKFN9 containing a HgaI site immediately after the MFl leader sequence is described in European patent application No. 0214826.

25 The ligation mixture was used to transform a competent E. coli strain ( $r^-$ ,  $m^+$ ) selecting for ampicillin resistance. Sequencing of a  $^{32}$ P-XbaI-EcoRI fragment (Maxam, A. and Gilbert, W. (1980) Methods Enzymol. 65, 499-560) showed that plasmids from the resulting colonies  
30 contained the correct DNA-sequence for aprotinin(3-58; 42 Ser) ..

One plasmid pKFN306 was selected for further use. The construction of plasmid pKFN306 is illustrated in fig. 2.

To introduce Ala in position 17 the following  
5 oligonucleotides were synthesized as described above:

Ia: CTGGTCCATGTAAAGCTGCTATCATCAGATACTTCTACAACGC

43-mer

10 IIa: CTTGGCGTTGTAGAAGTATCTGATGATAGCAGCTTTACATGGACCAGTGT

50-mer

The oligonucleotides were 5'-O-phosphorylated by treatment with ATP and  $T_4$ -kinase.

15 A duplex formed by annealing 5'-phosphorylated oligonucleotides Ia and IIa was ligated to the 352 bp EcoRI-PflMI fragment and the 3 kbp EcoRI-StyI fragment, both from pKFN306. pKFN306 encodes the S.cerevisiae mating factor  $\alpha 1$  signal-leader (1-85) fused to the synthetic  
20 aprotinin (3-58; 42 Ser) gene.

The ligation mixture was used to transform a competent E. coli strain ( $r^-$ ,  $m^+$ ) selecting for ampicillin resistance. Sequencing of a  $^{32}P$ -XbaI-EcoRI fragment (Maxam, A. and Gilbert, W. (1980) Methods Enzymol. 65, 499-560)  
25 showed that plasmids from the resulting colonies contained the correct DNA-sequence for aprotinin(3-58; 17Ala + 42Ser).

One plasmid, pKFN501 was selected for further use. The construction of plasmid pKFN501 is illustrated in  
30 fig. 3.

pKFN501 was cut with EcoRI and XbaI and the 0.5 kb fragment was ligated to the 9.5 kb NcoI-XbaI fragment from PMT636 and the 1.4 kb NcoI-EcoRI fragment from PMT636,

resulting in plasmid pKFN504, see fig. 3. Plasmid pMT636 was constructed from pMT608 after deletion of the LEU-2 gene and from pMT479, see fig. 4. pMT608 is described in European patent application No. 195691. pMT479 is described in European patent application No. 163529. pMT479 contains the Schizo. pombe TPI gene (POT), the S. cerevisiae triosephosphate isomerase promoter and terminator, TPI<sub>P</sub> and TPI<sub>T</sub> (Alber, T. and Kawasaki, G. (1982) J.Mol.Appl.Gen. 1, 419-434). Plasmid pKFN504 contains the following sequence:

100

TPI<sub>P</sub>-MFAI-signal-leader(1-85)-aprotinin(3-58;17Ala + 42Ser)-TPI<sub>T</sub> where MFAI is the S. cerevisiae mating factor alpha 1 coding sequence (Kurjan, J. and Herskowitz, I. (1982) Cell 30, 933-943), signal-leader(1-85) means that the sequence contains the first 85 amino acid residues of the MFAI signal-leader sequence and aprotinin(3-58; 17Ala + 42Ser) is the synthetic sequence encoding an aprotinin derivative lacking the first two amino acid residues at the N-terminus and having amino acid residues 17 and 42 replaced by an Ala and a Ser residue, respectively.

S. cerevisiae strain MT663 (E2-7B XE11-36 a/α, ΔtpiΔtpi, pep 4-3/pep 4-3) was grown on YPGaL (1% Bacto yeast extract, 2% Bacto peptone, 2% galactose, 1% lactate) to an optical density at 600nm of 0.6.

25

100 ml of culture was harvested by centrifugation, washed with 10 ml of water, recentrifuged and resuspended in 10 ml of (1.2 M sorbitol, 25 mM Na<sub>2</sub>EDTA pH = 8.0, 6.7 mg/ml dithiotreitol). The suspension was incubated at 30°C for 15 minutes, centrifuged and the cells resuspended in 10 ml of (1.2 M sorbitol, 10 mM Na<sub>2</sub>EDTA, 0.1 M sodium citrate pH = 5.8, 2 mg Novozym® 234). The suspension was incubated at 30°C for 30 minutes, the cells collected by centrifugation, washed in 10 ml of 1.2 M

30



sorbitol and 10 ml of CAS (1.2 M sorbitol, 10 mM  $\text{CaCl}_2$ , 10 mM Tris HCl (Tris = Tris(hydroxymethyl)-aminomethan) pH = 7.5) and resuspended in 2 ml of CAS. For transformation 0.1 ml of CAS-resuspended cells were mixed with approximately 1  
5  $\mu\text{g}$  of plasmid pKFN504 and left at room temperature for 15 minutes. 1 ml of (20% polyethylenglycol 4000, 10 mM  $\text{CaCl}_2$ , 10 mM Tris HCl, pH = 7.5) was added and the mixture left for further 30 minutes at room temperature. The mixture was centrifuged and the pellet resuspended in 0.1 ml of SOS  
10 (1.2 M sorbitol, 33% v/v YPD, 6.7 mM  $\text{CaCl}_2$ , 14  $\mu\text{g}/\text{ml}$  leucine) and incubated at 30°C for 2 hours. The suspension was then centrifuged and the pellet resuspended in 0.5 ml of 1.2 M sorbitol. 6 ml of top agar (the SC medium of Sherman et al., (Methods in Yeast Genetics, Cold Spring  
15 Harbor Laboratory, 1981) containing 1.2 M sorbitol plus 2.5% agar) at 52°C was added and the suspension poured on top of plates containing the same agar-solidified, sorbitol containing medium. Transformant colonies were picked after 3 days at 30°C, reisolated and used to start liquid  
20 cultures. One such transformant KFN396 was chosen for further characterization.

Yeast strain KFN396 was grown on YPD medium (1% yeast extract, 2% peptone (from Difco Laboratories), and 2% glucose). A 1 liter culture of the strain was shaken at  
25 30°C to an optical density of 600nm of 13. After centrifugation the supernatant was purified by FPLC ion exchange chromatography. The yeast supernatant was filtered through a 0.22  $\mu\text{m}$  Millex® GV filter unit and 1 ml was applied on a MonoS cation exchange column (0.5 x 5 cm)  
30 equilibrated with 20 mM Bicine, pH 8.7. After wash with equilibration buffer the column was eluted with a linear NaCl gradient (0-1 M) in equilibration buffer. Trypsin

inhibitor activity was quantified in the eluted fractions by spectrophotometric assay and furthermore by integration of absorption at 280 nm from

$$E_{280}^{1\%}(\text{aprotinin}) = 8.3$$

5

280

The yield was about 4.3 mg/liter of aprotinin(3-58; 17Ala + 42Ser).

For amino acid analysis the yeast supernatant (7 ml) was adjusted to pH 8.7 with 0.1 M NaOH and filtered (0.22  $\mu$ m). The effluent from a Q-Sepharose anion exchange column (11 x 4 cm) equilibrated with 20 mM Bicine, pH 8.7 was applied to a MonoS cation exchange column (0.5 x 5 cm). The cation exchange chromatography was carried out as described above. Concentration of the gradient eluted aprotinin(3-58) was made by rechromatography on MonoS and elution with steep NaCl-gradient. The collected fractions were further concentrated by vacuum centrifugation to about 100  $\mu$ l and applied to a RP-HPLC column (Vydac C4, 4.6 x 250 mm). Elution was carried out with CH<sub>3</sub>CN gradient in 0.1% TFA. The collected fractions were concentrated to about 100  $\mu$ l by vacuum centrifugation and samples were taken for amino acid analysis.

The amino acid analysis appears from the following Table 1. From this table it appears that the product has the expected amino acid composition.

Table 1

5	Amino acid	Theoretical	Aprotinin
			(3-58;17Ala + 42Ser) (found)
10	Asx	5	4.90
	Thr	3	2.95
	Ser	2	2.10
	Glx	3	3.01
	Pro	3	3.14
	Gly	6	5.93
	Ala	7	6.69
15	Cys	6	5.91
	Val	1	1.02
	Met	1	0.99
	Ile	2	2.00
	Leu	2	1.98
20	Tyr	4	3.73
	Phe	4	3.75
	Lys	4	4.29
	Arg	3	3.21
Total		56	55.60

25

Example 2Aprotinin(3-58;17Ala + 19Glu + 42Ser) (KFN399)

A synthetic gene encoding aprotinin(3-58; 17Ala  
 30 + 19Glu + 42Ser) was constructed as described in Example 1.  
 The following oligonucleotides Ib and IIb were used instead  
 of Ia and IIa:

Ib: CTGGTCCATGTAAAGCTGCTATCGAAAGATACTTCTACAACGC  
43-mer

IIb: CTTGGCGTTGTAGAAGTATCTTTTCGATAGCAGCTTTACATGGACCAGTGT  
5 50-mer

The pUC19 derived plasmid pKFN503 was constructed in a similar way as pKFN501.

100 By following the procedure of Example 1 a plasmid pKFN507 was obtained containing the following construction

TPI<sub>P</sub>-MfoI-signal-leader(1-85)-aprotinin(3-58;17Ala + 19Glu  
15 + 42Ser)-TPI<sub>T</sub>

where aprotinin(3-58; 17Ala + 19Glu + 42Ser) is the synthetic gene encoding an aprotinin derivative lacking the first two amino acid residues at the N-terminal and having  
20 the residues 17, 19 and 42 of native aprotinin replaced by an alanine, a glutamic acid and a serine residue, respectively.

Plasmid pKFN507 was transformed in yeast strain MT663 as described above and culturing of the transformed  
25 strain KFN399 gave about 10 mg/liter of aprotinin(3-58; 17Ala + 19Glu + 42Ser).

The amino acid analysis appears from the following Table 2 and confirms the expected amino acid composition.

Table 2

Amino Acid		Theoretical	Aprotinin
5			(3-58,17Ala + 19Glu + 42Ser)
			(found)
		<hr/>	
	Asx	5	4.95
	Thr	3	2.83
10	Ser	2	1.90
	Glx	4	4.08
	Pro	3	2.98
	Gly	6	5.98
	Ala	7	6.92
15	Cys	6	5.06
	Val	1	0.99
	Met	1	0.86
	Ile	1	0.99
	Leu	2	1.99
20	Tyr	4	3.77
	Phe	4	3.89
	Lys	4	4.07
	Arg	3	3.06
		<hr/>	
	Total	56	54.36
		<hr/>	

25

Example 3Aprotinin(3-58;15Arg + 17Ala + 42Ser)(KFN773).

A synthetic gene encoding aprotinin(3-58; 15Arg + 17Ala + 42Ser) was constructed as described in Example 1.

- 5 The following oligonucleotides Ic and IIc were used instead of Ia and IIa:

Ic: CTGGTCCATGTAGAGCTGCTATCATCAGATACTTCTACAACGC

10 43-mer

IIc: CTTGGCGTTGTAGAAGTATCTGATGATAGCAGCTCTACATGGACCAGTGT

50-mer

- 15 The pUC19 derived plasmid pKFN777 was constructed in a similar way as pKFN501.

- By following the procedure of Example 1 a plasmid pKFN807 was obtained containing the following  
20 construction

TPI<sub>P</sub>-M<sub>F</sub>ol-signal-leader(1-85)-aprotinin(3-58;15Arg + 17Ala + 42Ser)-TPI<sub>T</sub>

- 25 where aprotinin(3-58; 15Arg + 17Ala + 42Ser) is the synthetic gene encoding an aprotinin derivative lacking the first two amino acid residues at the N-terminal and having the residues 15, 17 and 42 of native aprotinin replaced by an arginine, an alanine and a serine residue, respectively.

- 30 Plasmid pKFN807 was transformed in yeast strain MT663 as described above and culturing of the transformed strain KFN773 gave about 8.5 mg/liter of aprotinin(3-58; 15Arg + 17Ala + 42Ser).

The amino acid analysis is shown in Table 3 and confirms the expected amino acid composition.

5

Table 3

	Amino Acid	Theoretical	Aprotinin
			(3-58,15Arg + 17Ala + 42Ser) (found)
10			
	Asx	5	4.95
	Thr	3	2.85
	Ser	2	1.81
15	Glx	3	3.01
	Pro	3	3.05
	Gly	6	5.92
	Ala	7	6.91
	Cys	6	5.31
20	Val	1	1.02
	Met	1	0.73
	Ile	2	1.41
	Leu	2	1.99
	Tyr	4	3.80
25	Phe	4	3.94
	Lys	3	2.97
	Arg	4	4.24
	Total	56	53.91

30

The slightly lowered content of Ile compared with the theoretical value can most probably be ascribed to incomplete hydrolysis of Ile(18)-Ile(19). This is well known in the art.

Example 4

Inhibition of Serine Proteases from Plasma by Aprotinin(3-58; 17Ala + 42Ser) (KFN396) and Aprotinin(3-58; 17Ala + 19Glu + 42Ser) (KFN399), Aprotinin(3-58;15Arg + 42Ser)(KFN772) and Aprotinin(3-58; 15Arg + 17Ala + 42Ser)(KFN773).

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Aprotinin(3-58; 17Ala + 42Ser)(KFN396),  
10 aprotinin(3-58; 17Ala + 19Glu + 42Ser)(KFN399) and  
aprotinin (3-58; 15Arg + 17Ala + 42Ser)(KFN773) were  
purified as described above. As native, bovine pancreatic  
aprotinin(1-58) batch B 5029-65 (67000 KIU/mg) from NOVO  
(Bagsvaerd, Denmark) was used. The concentration was  
15 calculated using  $E_{280\text{nm}} = 8.3$  and  $M_r = 6500$ . Human plasma  
kallikrein was obtained from Sigma (St. Louis, MO), bovine  
factor Xa was purified according to (H. Nobukazu et al.  
J.Biochem. 97 (1985)1347-1355), human factor IIa (thrombin)  
was a gift from Dr. W. Lawson (New York State Dept. of  
20 Health, Albany, N.Y.), recombinant human factor VIIa was  
from NOVO (Bagsvaerd, Denmark) and recombinant human  
protein Ca was from ZymoGenetics, Inc. (Seattle, WA).  
Substrate S 2302 (H-D-Pro-Phe-Arg-p-nitroanilide) substrate  
S2238 (H-D-Phe-Pip-Arg-p-nitroanilide) and substrate S2366  
25 (Glu-Pro-Arg-p-nitroanilide) were from Kabi (Stockholm,  
Sweden). Substrate FXa-1 (methoxycarbonyl DCH-Gly-Arg-p-  
nitroanilide) was from NycoMed (Oslo, Norway). The  
experiments were performed in 100 mM NaCl, 50 mM Tris-HCl  
0.01 % Tween80, pH 7.4 at 25°C.

30 Human plasma kallikrein (3 nM) was incubated  
with aprotinin (0-20 nM) for 30 minutes in a micro-titer  
well. Substrate S 2302 (0.6 mM) was added to a final volume  
of 300  $\mu$ l, and the rate of nitroaniline generation was



measured at 405 nm by means of a Micro ELISA® Autoreader MR 580 from Dynatech Laboratories. The rate is proportional to the concentration of free enzyme. The inhibition of plasma kallikrein by native aprotinin and the four analogues KFN 396, KFN 399, KFN 772 and KFN 773 is shown in fig. 5A and 5B. With native aprotinin a moderate inhibition was observed. The inhibition was strongly increased by analogues KFN 396 and KFN 399 containing Ala in position 17 (Fig. 5A).

10 A further increase of the inhibition was obtained with Arg in position 15 (KFN 772); and the strongest inhibition was observed with the analogue (KFN 773) with substitution of both position 17 (Ala) and position 15 (Arg) (Fig. 5B).

15 The analogues were also tested for inhibition of the amidolytic activity of the serine proteases: bovine factor Xa, human factor IIa, human recombinant factor VIIa and human recombinant protein Ca. The experiments were performed essentially as described for plasma kallikrein only appropriate substrates were used. Finally the analogues were analysed for an effect on the coagulation factors of human plasma by means of two clotting tests. These tests, the prothrombin time (PTT) and the activated thromboplastin time (APTT) were performed with General

25 Diagnostics® reagents from Organon (Durham, NC) according to the directions given by the manufacturer. The results of the inhibition experiments are summarized in Table 4 which describes the inhibition profile of the four aprotinin analogues. KFN 773 is characterized by an extraordinarily strong inhibition of human plasma kallikrein, which is ten

30 fold stronger than that of the Arg 15 analogue (KFN 772). A reverse effect is observed with activated protein C. In

this case the relatively strong inhibition obtained by substitution of Lys 15 to Arg is weakened by further substitution of Arg 17 to Ala.

5 Table 4. Inhibition profile of aprotinin analogues

100	Product	$K_i^{*})$ (nM); Amidolytic activity of serine proteases§				Clot assays		
		Plasma kallikrein	FIIa	FVIIa	FXa	Prot.Ca	PTT	APTT
	Native Aprotinin	180	-	-	-	400	-	-
15	KFN 396	12	-	-	-		-	-
	KFN 399	12						
	KFN 772	1	-	-	1800	10	-	+
	KFN 773	0.1	-	-	150	100	-	+

20

- No inhibition at 1.0  $\mu$ M aprotinin analogue

+ Prolonged clotting time at 1.0  $\mu$ M aprotinin analogue

25 \*) Inhibition constants estimated according to the graphical Dixon method (M. Dixon, Biochem. J. 129 (1972)197-202)

§ Substrates: Plasma kallikrein: S2302; FIIa: S2238; FVIIa: Substrate FXa-1; FXa: Substrate FXa-1; Prot. Ca: S2366.

30

5

## CLAIMS

10:

15

1. Aprotinin analogues having the formula  
X<sub>1</sub>-Asp-Phe-Cys-Leu-Glu-Pro-Pro-Tyr-Thr-X<sub>2</sub>-X<sub>3</sub>-X<sub>4</sub>-X<sub>5</sub>-X<sub>6</sub>-X<sub>7</sub>-  
X<sub>8</sub>-X<sub>9</sub>-Arg-Tyr-Phe-Tyr-Asn-Ala-Lys-Ala-Gly-Leu-Cys-Gln-Thr-  
Phe-Val-Tyr-Gly-Gly-X<sub>10</sub>-Arg-Ala-X<sub>11</sub>-X<sub>12</sub>-Asn-Asn-Phe-Lys-  
20 Ser-Ala-Glu-Asp-Cys-Met-Arg-Thr-Cys-Gly-Gly-Ala in which X<sub>1</sub>  
means Arg-Pro, Pro or hydrogen, X<sub>4</sub> and X<sub>10</sub> are both Cys or  
both Ala or both another naturally occurring amino acid  
residue, X<sub>2</sub> and X<sub>3</sub> are independently any naturally  
occurring amino acid residue; X<sub>5</sub> is Lys, Arg, Val, Thr, Ile,  
25 Leu, Phe, Gly, Ser, Met, Trp, Tyr or Ala with the proviso  
that when X<sub>5</sub> is different from Lys then at least one of X<sub>2</sub>  
to X<sub>4</sub> and X<sub>6</sub> to X<sub>9</sub> is different from the amino acid residue  
at the corresponding position in native aprotinin; X<sub>6</sub> is  
Ala or Gly; X<sub>7</sub> is any naturally occurring amino acid  
30 residue; X<sub>8</sub> is Ile, Leu, Met, Val or Phe; X<sub>9</sub> is any  
naturally occurring amino acid residue; X<sub>11</sub> is any naturally  
occurring amino acid residue; X<sub>12</sub> is Lys, Arg or Ser at

least one of the amino acid residues  $X_2$  to  $X_9$  being different from the corresponding amino acid residue in native aprotinin.

2. Aprotinin analogues according to Claim 1,  
5 wherein  $X_1$  is Arg-Pro, Pro or hydrogen, preferably Arg-Pro or hydrogen, most preferred hydrogen.

3. Aprotinin analogues according to Claim 1 or 2, wherein  $X_2$  is Gly.

4. Aprotinin analogues according to any of the  
10 preceding claims, wherein  $X_3$  is Pro.

5. Aprotinin analogues according to any of the preceding claims, wherein  $X_4$  and  $X_{10}$  are Cys.

6. Aprotinin analogues according to any of the preceding claims wherein  $X_5$  is Lys, Arg, Val, Thr, Ile,  
15 Leu, Phe, Gly, Ser, Met, Trp, Tyr or Ala, preferably Lys or Arg with the proviso that when  $X_5$  is different from Lys, then at least one of  $X_2$  to  $X_4$  and  $X_6$  to  $X_9$ , preferably  $X_6$  to  $X_9$ , is different from the amino acid residue at the corresponding position in native aprotinin.

20 7. Aprotinin analogues according to any of the preceding claims, wherein  $X_6$  is Ala or Gly, preferably Ala.

8. Aprotinin analogues according to any of the preceding claims, wherein  $X_7$  is Ala or Gly, preferably Ala.

9. Aprotinin analogues according to any of the  
25 preceding claims wherein  $X_8$  is Ile, Leu, Met, Val or Phe, preferably Ile.

10. Aprotinin analogues according to any of the preceding claims wherein  $X_9$  is Ile or Glu, preferably Ile.

11. Aprotinin analogues according to any of the  
30 preceding claims wherein  $X_{11}$  is Lys or Arg, preferably Lys.

12. Aprotinin analogues according to any of the preceding claims wherein  $X_{12}$  is Lys, Arg or Ser, preferably Arg or Ser.

13. Aprotinin analogue according to claim 1 having the following structure:

Arg-Pro-Asp-Phe-Cys-Leu-Glu-Pro-Pro-Tyr-Thr-Gly-Pro-Cys-  
Lys-Ala-Ala-Ile-Glu-Arg-Tyr-Phe-Tyr-Asn-Ala-Lys-Ala-Gly-  
5 Leu-Cys-Gln-Thr-Phe-Val-Tyr-Gly-Gly-Cys-Arg-Ala-Lys-Arg-  
Asn-Asn-Phe-Lys-Ser-Ala-Glu-Asp-Cys-Met-Arg-Thr-Cys-Gly-  
Gly-Ala.

14. Aprotinin analogue according to claim 1 having the following structure:

10 Asp-Phe-Cys-Leu-Glu-Pro-Pro-Tyr-Thr-Gly-Pro-Cys-Lys-Ala-  
Ala-Ile-Glu-Arg-Tyr-Phe-Tyr-Asn-Ala-Lys-Ala-Gly-Leu-Cys-  
Gln-Thr-Phe-Val-Tyr-Gly-Gly-Cys-Arg-Ala-Lys-Arg-Asn-Asn-  
Phe-Lys-Ser-Ala-Glu-Asp-Cys-Met-Arg-Thr-Cys-Gly-Gly-Ala.

15 15. Aprotinin analogue according to claim 1 having the following structure:

Arg-Pro-Asp-Phe-Cys-Leu-Glu-Pro-Pro-Tyr-Thr-Gly-Pro-Cys-  
Lys-Ala-Ala-Ile-Glu-Arg-Tyr-Phe-Tyr-Asn-Ala-Lys-Ala-Gly-  
Leu-Cys-Gln-Thr-Phe-Val-Tyr-Gly-Gly-Cys-Arg-Ala-Lys-Ser-  
Asn-Asn-Phe-Lys-Ser-Ala-Glu-Asp-Cys-Met-Arg-Thr-Cys-Gly-  
20 Gly-Ala.

16. Aprotinin analogue according to claim 1 having the following structure:

Asp-Phe-Cys-Leu-Glu-Pro-Pro-Tyr-Thr-Gly-Pro-Cys-Lys-Ala-  
Ala-Ile-Glu-Arg-Tyr-Phe-Tyr-Asn-Ala-Lys-Ala-Gly-Leu-Cys-  
25 Gln-Thr-Phe-Val-Tyr-Gly-Gly-Cys-Arg-Ala-Lys-Ser-Asn-Asn-  
Phe-Lys-Ser-Ala-Glu-Asp-Cys-Met-Arg-Thr-Cys-Gly-Gly-Ala.

17. Aprotinin analogue according to claim 1 having the following structure:

Arg-Pro-Asp-Phe-Cys-Leu-Glu-Pro-Pro-Tyr-Thr-Gly-Pro-Cys-  
30 Lys-Ala-Ala-Ile-Ile-Arg-Tyr-Phe-Tyr-Asn-Ala-Lys-Ala-Gly-  
Leu-Cys-Gln-Thr-Phe-Val-Tyr-Gly-Gly-Cys-Arg-Ala-Lys-Arg-  
Asn-Asn-Phe-Lys-Ser-Ala-Glu-Asp-Cys-Met-Arg-Thr-Cys-Gly-  
Gly-Ala.

18. Aprotinin analogue according to claim 1  
having the following structure:

Asp-Phe-Cys-Leu-Glu-Pro-Pro-Tyr-Thr-Gly-Pro-Cys-Lys-Ala-  
Ala-Ile-Ile-Arg-Tyr-Phe-Tyr-Asn-Ala-Lys-Ala-Gly-Leu-Cys-  
5 Gln-Thr-Phe-Val-Tyr-Gly-Gly-Cys-Arg-Ala-Lys-Arg-Asn-Asn-  
Phe-Lys-Ser-Ala-Glu-Asp-Cys-Met-Arg-Thr-Cys-Gly-Gly-Ala.

19. Aprotinin analogue according to claim 1  
having the following structure:

Asp-Phe-Cys-Leu-Glu-Pro-Pro-Tyr-Thr-Gly-Pro-Cys-Lys-Ala-  
100 ~~Ala-Ile-Ile-Arg-Tyr-Phe-Tyr-Asn-Ala-Lys-Ala-Gly-Leu-Cys-~~  
Gln-Thr-Phe-Val-Tyr-Gly-Gly-Cys-Arg-Ala-Lys-Ser-Asn-Asn-  
Phe-Lys-Ser-Ala-Glu-Asp-Cys-Met-Arg-Thr-Cys-Gly-Gly-Ala.

20. Aprotinin analogue according to claim 1  
having the following structure:

15 Arg-Pro-Asp-Phe-Cys-Leu-Glu-Pro-Pro-Tyr-Thr-Gly-Pro-Cys-  
Lys-Ala-Ala-Ile-Ile-Arg-Tyr-Phe-Tyr-Asn-Ala-Lys-Ala-Gly-  
Leu-Cys-Gln-Thr-Phe-Val-Tyr-Gly-Gly-Cys-Arg-Ala-Lys-Ser-  
Asn-Asn-Phe-Lys-Ser-Ala-Glu-Asp-Cys-Met-Arg-Thr-Cys-Gly-  
Gly-Ala.

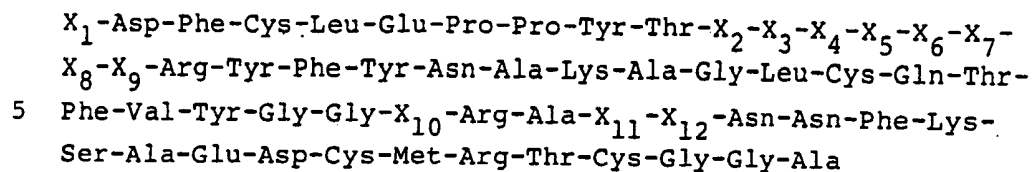
20 21. Aprotinin analogue according to claim 1  
having the following structure:

Asp-Phe-Cys-Leu-Glu-Pro-Pro-Tyr-Thr-Gly-Pro-Cys-Arg-Ala-  
Ala-Ile-Ile-Arg-Tyr-Phe-Tyr-Asn-Ala-Lys-Ala-Gly-Leu-Cys-  
Gln-Thr-Phe-Val-Tyr-Gly-Gly-Cys-Arg-Ala-Lys-Ser-Asn-Asn-  
25 Phe-Lys-Ser-Ala-Glu-Asp-Cys-Met-Arg-Thr-Cys-Gly-Gly-Ala.

22. Aprotinin analogue according to claim 1  
having the following structure:

Arg-Pro-Asp-Phe-Cys-Leu-Glu-Pro-Pro-Tyr-Thr-Gly-Pro-Cys-  
Arg-Ala-Ala-Ile-Ile-Arg-Tyr-Phe-Tyr-Asn-Ala-Lys-Ala-Gly-  
30 Leu-Cys-Gln-Thr-Phe-Val-Tyr-Gly-Gly-Cys-Arg-Ala-Lys-Ser-  
Asn-Asn-Phe-Lys-Ser-Ala-Glu-Asp-Cys-Met-Arg-Thr-Cys-Gly-  
Gly-Ala.

23. A process for producing aprotinin analogues according to the general formula (I):



(I)

wherein  $X_1$  is Arg-Pro, Pro or hydrogen;  $X_2$  is any naturally  
 10 occurring amino acid residue, preferably Gly;  $X_3$  is any  
 naturally occurring amino acid residue, preferably Pro;  $X_4$   
 and  $X_{10}$  are any naturally occurring amino acid residues,  
 preferably they are both Ala or both Cys;  $X_5$  is Lys, Arg,  
 Val, Thr, Ile, Leu, Phe, Gly, Ser, Met, Trp, Tyr or Ala,  
 15 preferably Lys or Arg, with the proviso, that when  $X_5$  is  
 different from Lys, then at least one of  $X_2$  to  $X_4$  and  $X_6$  to  
 $X_9$  is different from the amino acid residue at the  
 corresponding position in native aprotinin;  $X_6$  is Ala or  
 Gly, preferably Ala;  $X_7$  is any naturally occurring amino  
 20 acid residue, preferably Ala or Gly;  $X_8$  is Ile, Leu, Met,  
 Val or Phe, preferably Ile;  $X_9$  is any naturally occurring  
 amino acid residue, preferably Ile or Glu;  $X_{11}$  is any  
 naturally occurring amino acid residue, preferably Lys or  
 Arg and  $X_{12}$  is Lys, Arg or Ser, preferably Arg or Ser, at  
 25 least one of the amino acid residues  $X_2$  to  $X_9$ , preferably  
 $X_5$  to  $X_9$ , more preferred  $X_6$  to  $X_9$  being different from the  
 corresponding amino acid residue in native protein in yeast  
 comprising culturing a yeast strain containing a replicable  
 expression vector, the vector comprising a gene coding for  
 30 the aprotinin analogue and DNA sequences that allow for the  
 expression of the aprotinin analogue in a suitable nutrient  
 medium and recovering the expressed aprotinin analogues.

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3 5 10 15  
AspPheCysLeuGluProProTyrThrGlyProCysLysAlaArgIle  
AAAGAGATTCTGTTGGAACCTCCATACTGGTCCATGTAAAGCTAGAATC  
CTAAAGACAAACCTTGGAGGTATGTGACCAGGTACATTTTCGATCTTAG  
PflMI

20 25 30 35  
IleArgTyrPheTyrAsnAlaLysAlaGlyLeuCysGlnThrPheValTyrGly  
ATCAGATACTTCTACAACGCCAAGGCTGGTTTGTGTCAAACCTTTCGTTTACGGT  
TAGTCTATGAAGATGTTGCGGTTCCGACCAAACACAGTTTGAAAGCAAATGCCA  
StyI

40 45 50  
GlyCysArgAlaLysSerAsnAsnPheLysSerAlaGluAspCysMetArgThr  
GGCTGCAGAGCTAAGTCCAACAACCTTCAAGTCTGCTGAAGACTGCATGAGAACT  
CCGACGTCTCGATTTCAGGTTGTTGAAGTTCAGACGACTTCTGACGTACTCTTGA  
PstI

55 58  
CysGlyGlyAlaStop  
TGTGGTGGTGCCTAAT  
ACACCACCACGGATTAGATC  
XbaI

FIG. 1



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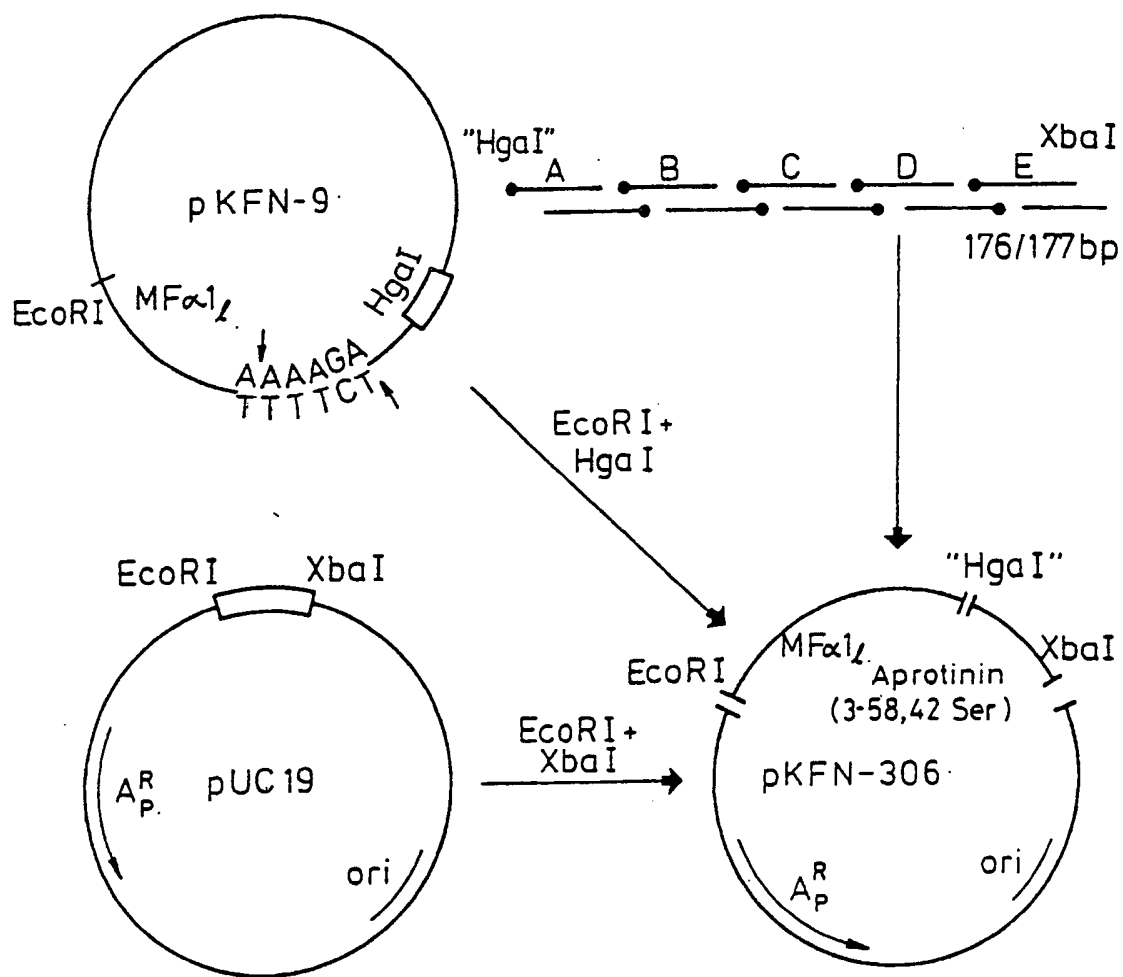
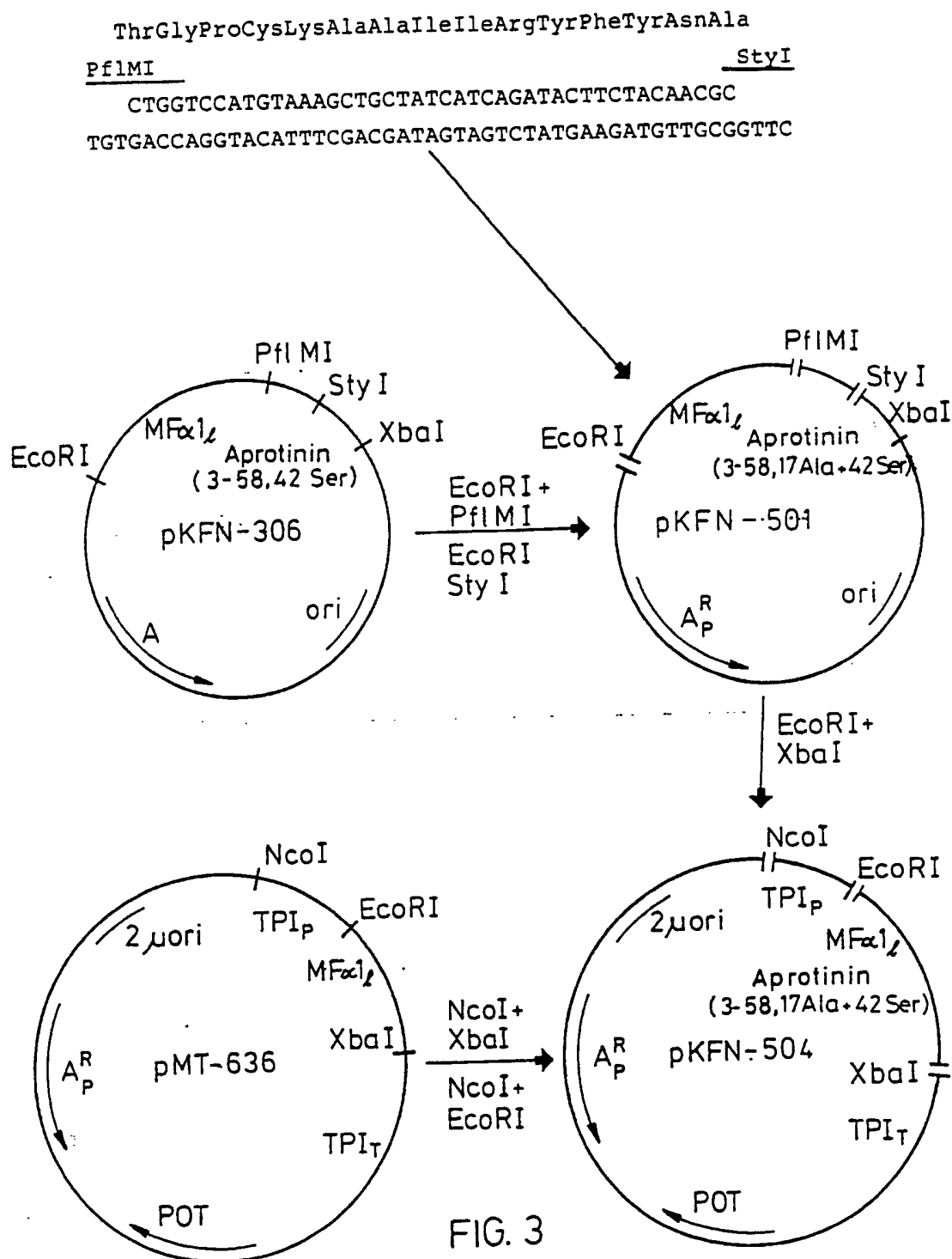


FIG. 2

3/5



SUBSTITUTE

4 / 5

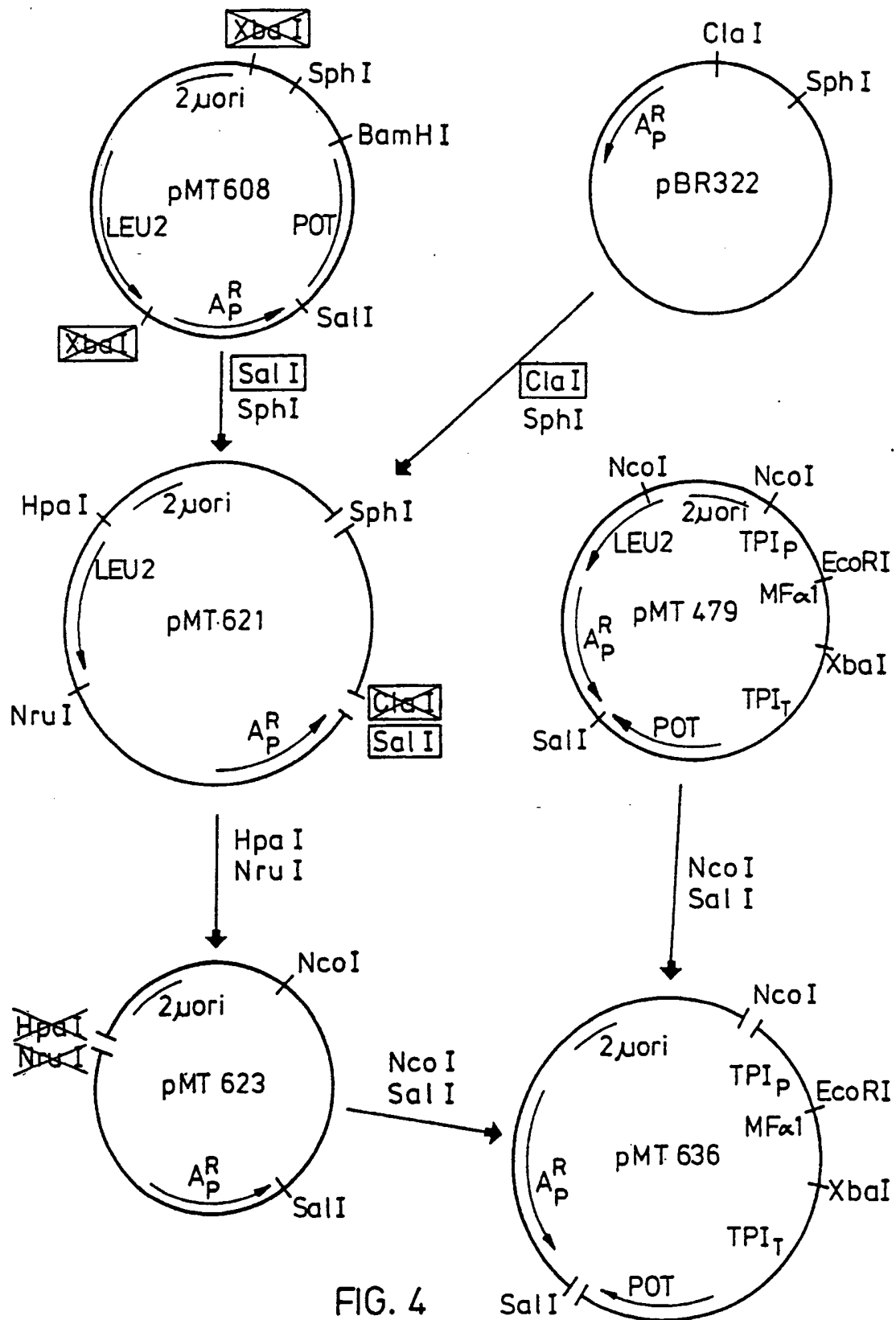


FIG. 4

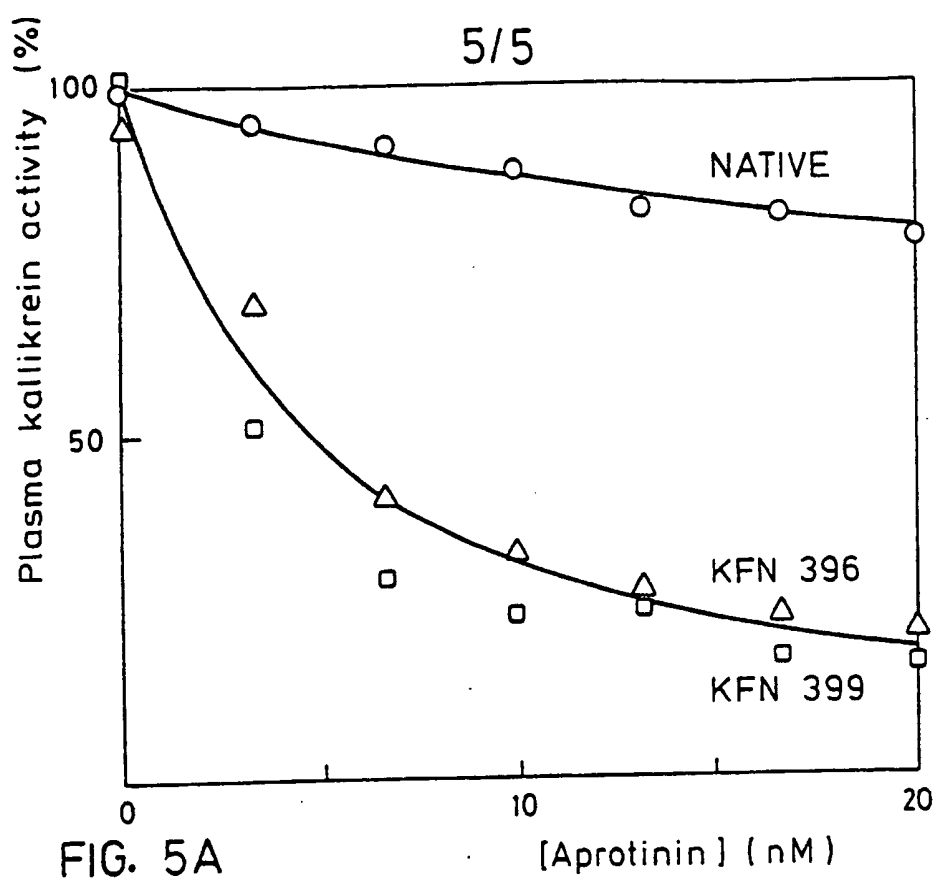


FIG. 5A

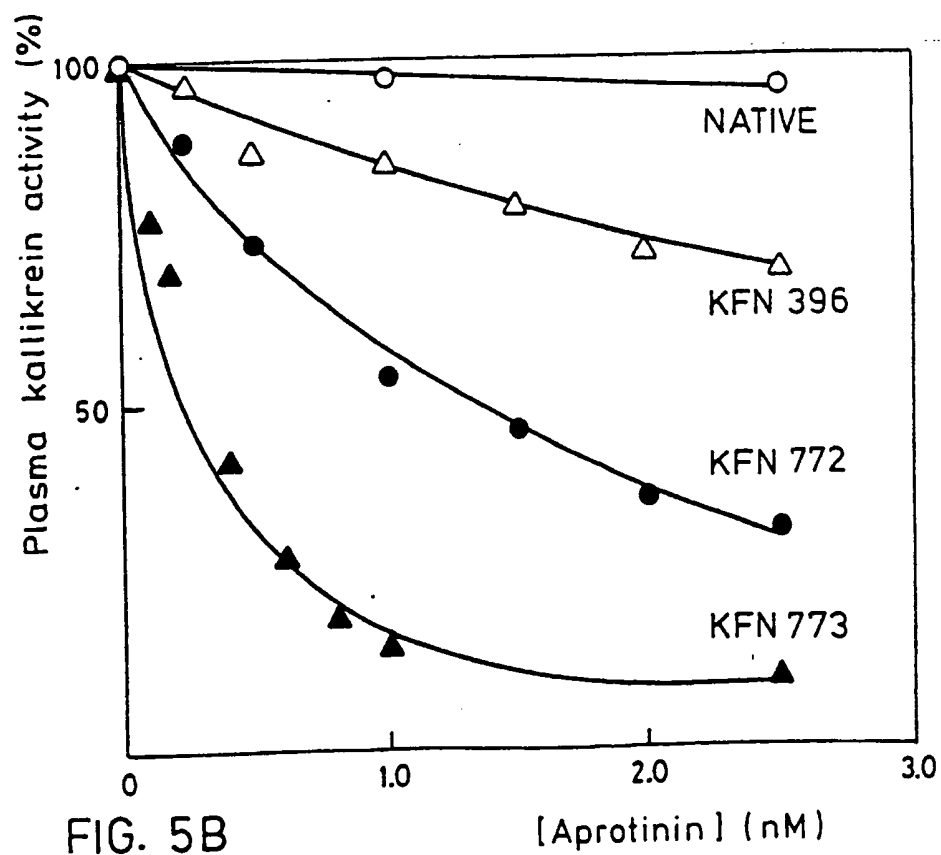


FIG. 5B

# INTERNATIONAL SEARCH REPORT

International Application No. PCT/DK89/00096

<b>I. CLASSIFICATION OF SUBJECT MATTER</b> (if several classification symbols apply, indicate all) *		
According to International Patent Classification (IPC) or to both National Classification and IPC 4		
C 07 K 7/10, C 12 N 15/00		
<b>II. FIELDS SEARCHED</b>		
Minimum Documentation Searched 7		
Classification System	Classification Symbols	
IPC 4	C07C 103/52; C07K 7/10	
US C1	260:112.5, 424:177; 514:2, 12; 530:324	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched *		
SE, NO, DK, FI classes as above		
<b>III. DOCUMENTS CONSIDERED TO BE RELEVANT *</b>		
Category *	Citation of Document, 11 with indication, where appropriate, of the relevant passages 12	Relevant to Claim No. 13
Y	EP, A2, 0 132 732 (BAYER AG) 13 February 1985 JP, 60056999 US, 4595674 AU, 560584 CA, 1238000 DE, 3471940 DE, 3339693	1-12, 22
Y	EP, A2, 0 238 993 (BAYER AG) 30 September 1987 GB, 2188322 JP, 63007794	1-12, 22 23
Y	EP, A2, 0 244 627 (BAYER AG) 11 November 1987 GB, 2188933 JP, 63012299	1-12, 23
P, X	EP, A2, 0 297 362 (BAYER AG) 4 January 1989 DE, 3724570	1-23
<p>* Special categories of cited documents: 10</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"A" document member of the same patent family</p>		
<b>IV. CERTIFICATION</b>		
Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report	
1989-07-14	1989-07-28	
International Searching Authority	Signature of Authorized Officer	
Swedish Patent Office	Elisabeth Carlborg	

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category *	Citation of Document, with indication, where appropriate, of the relevant passages	Relevant to Claim No
X	Biol. Chem. Hoppe-Seyler	1-4
Y	Vol. 367, pp. 1167-1176, November 1986	5-12
	Eugen Schnabel et al	
	"[Ala <sub>2</sub> <sup>14,38</sup> ]Aprotinin: Preparation by Partial Desulphurization of Aprotinin by Means of Raney Nickel and Comparison with Other Aprotinin Derivatives".	